

## METHODS FOR STABILIZING LYOPHILIZED BLOOD PROTEINS

### FIELD OF THE INVENTION

The present invention relates to methods for stabilizing lyophilized blood proteins using hydroxypropyl- $\alpha$ -cyclodextrin.

### BACKGROUND OF THE INVENTION

Fibrinogen is an important blood protein. Fibrinogen-containing solutions can be infused intravenously as replacement therapy for afibrogenemic patients. They are also a component of fibrin glue (FG) preparations. FG contains two components, fibrinogen and thrombin, which, when mixed together, form a "glue" for wound closure or for producing hemostasis at an injury site. Each component is supplied as a freeze-dried powder that must be reconstituted with diluent prior to use. After reconstitution, each component is delivered by an application device to the wound site, at which time the components are mixed and clotting (glue formation) occurs. However, most freeze-dried preparations of fibrinogen require relatively long periods of time for rehydration, and may denature or aggregate to form insoluble particles. Thus a need exists for an improved method for stabilizing fibrinogen to minimize potential denaturation and aggregation of the protein and reduce the rehydration time.

### SUMMARY OF THE INVENTION

The present invention provides a process for stabilizing lyophilized blood proteins. In one embodiment, the invention is directed to a method for stabilizing lyophilized blood proteins, particularly lyophilized fibrinogen. The method comprises providing an aqueous solution of a blood protein. Hydroxypropyl- $\alpha$ -cyclodextrin is added to the solution in an amount sufficient to form a complex with at least a portion, and preferably all, of the blood protein. The solution is lyophilized to provide a dry blood protein/hydroxypropyl- $\alpha$ -cyclodextrin complex. The dry blood protein/hydroxypropyl- $\alpha$ -cyclodextrin complex may then be reconstituted to provide a solution of the blood protein, which can be administered to a patient.

It has been discovered that the stabilization of blood protein with hydroxypropyl- $\alpha$ -cyclodextrin prior to lyophilization can reduce denaturation of the protein during dry heat viral inactivation. Additionally, the reconstitution time for the lyophilized blood protein stabilized in accordance with practice of the present invention is substantially reduced.

# DETAILED DESCRIPTION

The present invention is directed to a method that incorporates the use of hydroxypropyl- $\alpha$ -cyclodextrin (HP $\alpha$ CD) to stabilize lyophilized proteins, particularly fibrinogen, and to enhance reconstitution of these proteins. The method comprises providing an aqueous solution of a blood protein. HP $\alpha$ CD is added to the solution in an amount sufficient to form a complex with at least part, and preferably all, of the blood protein. The complex is lyophilized to provide a dry blood protein/HP $\alpha$ CD complex. The dry blood protein/HP $\alpha$ CD complex may then be reconstituted to provide a solution of the blood protein, which can be administered to a patient.

Blood proteins with which the present process may be used include, but are not limited to, albumin, Factor II, Factor VII, Factor VIII, Factor IX, Factors X and X<sub>a</sub>, fibrinogen, antithrombin III, transferrin, haptoglobin, gamma globulins, fibronectin, protein C, protein S and thrombin.

Cyclodextrins are homologous oligosaccharides that are obtained from starch by the action of enzymes from *Bacillus macetans*.  $\alpha$ -Cyclodextrin is a cyclic molecule containing six  $\alpha$ -D-glucopyranose units linked together at the 1,4 positions, as in amylose. This cyclic structure may also be referred to as a torus. HP $\alpha$ CD is commercially available from Cerestar USA, Inc. (Hammond, Indiana) or Pfanstiehl (Waukegan, Illinois).

The HP $\alpha$ CD may be added to an aqueous solution containing the blood protein before lyophilization at any suitable point in the purification process. Preferably, the HP $\alpha$ CD is added to an aqueous solution of the blood protein after all purification steps have been completed to prevent the HP $\alpha$ CD from forming a complex with impurities, which makes removal of the impurities more difficult.

If desired, the blood protein can be subjected to one or more viral inactivation steps prior to lyophilization, and preferably prior to complexing with the HP $\alpha$ CD. After lyophilization, preferably the blood protein is heated to a temperature and for a time sufficient to inactivate any viral contaminants. Preferably the complex is heated to a temperature of at least about 60°C, more preferably to at least about 80°C, still more preferably at least about 100°C, for a time of at least about 10 hours at 80°C or at least about 1 hour at 100°C, and more preferably at least about 72 hours at 80°C or at least about 3 hours at 100°C.

The blood protein can be subjected to a solvent detergent viral inactivation process instead of or in addition to viral inactivation by heat. Suitable solvent detergent viral inactivation processes are described in U.S. Patents Nos. 4,540,573, and 4,764,369, the entire disclosures of which are incorporated herein by reference.

Preferably the HP $\alpha$ CD is added in an amount sufficient to assure the formation of a complex with all of the desired blood protein. More preferably the HP $\alpha$ CD is added in an amount such that the aqueous solution has a HP $\alpha$ CD concentration of at least about 0.5% weight per volume (wt/vol.), preferably from about 0.5% to about 15% wt/vol., and more preferably from about 1% to about 12% wt/vol. More particularly, when the blood protein is fibrinogen, preferably the HP $\alpha$ CD is added in an amount such that the aqueous solution has a HP $\alpha$ CD concentration of at least about 0.5% wt/vol., preferably from about 0.5% to about 4% wt/vol., and more preferably from about 1% to about 2.5% wt/vol.

It has been found that the presence of HP $\alpha$ CD substantially decreases the reconstitution time of the lyophilized blood protein. The time for reconstituting the lyophilized protein/hydroxypropyl- $\alpha$ -cyclodextrin complex, compared to the time for reconstituting a similar protein solution not containing hydroxypropyl- $\alpha$ -cyclodextrin, is preferably decreased by at least about 50%, more preferably by at least about 75%, still more preferably by at least about 90%, and even more preferably by at least about 95%.

If desired, an additional stabilizing agent can be included with the HP $\alpha$ CD to further reduce the reconstitution time. Examples of such agents include lysine and polysorbate 80 (Tween 80).

### Example 1

Fibrinogen is manufactured from pooled cryo-poor and/or PTC-poor human plasma maintained at a temperature of  $1.5 \pm 1.5^\circ\text{C}$ . The pH is adjusted to  $7.0 \pm 0.2$  with either 1 M sodium bicarbonate or pH 4.0 acetate buffer. Sufficient cold SD3A ethanol is added to bring the plasma to a final alcohol concentration of 8%. During the alcohol addition, the temperature is gradually lowered to  $-2 \pm 4^\circ\text{C}$ . The precipitate that forms (Fraction I precipitate) is removed by centrifugation at  $-2 \pm 1^\circ\text{C}$ .

The Fraction I precipitate is extracted with about  $9 \pm 2$  kg of extraction buffer ( $0.40 \pm 0.15$  M 6-amino-n-hexanoic acid;  $0.05 \pm 0.01$  M sodium citrate;  $0.08 \pm 0.02$  M sodium chloride;  $7 \pm 4$  units/mL heparin; pH  $6.4 \pm 0.3$ ) per kg of Fraction I precipitate at pH  $6.4 \pm 0.3$ . Reconstitution of the Fraction I precipitate is performed at  $30 \pm 4^\circ\text{C}$  and yields Fraction I Solution. The pH of Fraction I Solution is adjusted to  $6.6 \pm 0.3$  if necessary. The extracted Fraction I solution is clarified by centrifugation and/or filtration at  $28 \pm 6^\circ\text{C}$  to produce Fraction I Filtrate.

Each kilogram of Fraction I Filtrate is mixed with  $0.11 \pm 0.03$  kg of Solvent Detergent Solution ( $3 \pm 0.5\%$  tri-n-butyl phosphate;  $10 \pm 1\%$  polysorbate 80; water for injection) to a final

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concentration of  $0.30 \pm 0.1\%$  tri-n-butyl phosphate and  $1 \pm 0.3\%$  polysorbate 80, and the pH of the mixture is adjusted to  $6.6 \pm 0.3$ . The solution is mixed for 1 hour at  $27 \pm 3^\circ\text{C}$  and transferred for further processing to a virally controlled area. Mixing is continued in the virally controlled area for an additional  $6 \pm 1$  hours at  $27 \pm 3^\circ\text{C}$ . The pH is adjusted as necessary to  $6.6 \pm 0.3$  during incubation.

The solution is cooled to  $23 \pm 4^\circ\text{C}$ , and the pH is adjusted to  $6.8 \pm 0.3$  with 1 N sodium hydroxide or 1 N acetic acid. The pH adjusted solution is cooled to  $9 \pm 3^\circ\text{C}$  with constant mixing, and solid glycine is added ( $135 \pm 25$  g per kg of pH adjusted solution). Mixing is continued for not less than 30 minutes at 3 to  $11^\circ\text{C}$  to obtain a First Glycine Precipitate. The First Glycine Precipitate is removed from the suspension by centrifugation or filtration and may be held frozen prior to further processing.

The First Glycine Precipitate is solubilized in approximately  $9 \pm 2$  kg of citrate saline buffer ( $0.02 \pm 0.005\text{M}$  sodium citrate;  $0.12 \pm 0.03$  M sodium chloride; pH  $7.7 \pm 0.5$ ) per kg of precipitate by mixing for at least 30 minutes at  $30 \pm 4^\circ\text{C}$ . The First Glycine Precipitate suspension is cooled to  $23 \pm 4^\circ\text{C}$ , and the pH is adjusted to  $6.8 \pm 0.3$  with 1 N acetic acid or 1 N sodium hydroxide. The adjusted solution is cooled to  $9 \pm 3^\circ\text{C}$  with constant mixing.

Solid glycine is added to the pH adjusted solution ( $128 \pm 20$  g per kg of adjusted solution) with vigorous mixing of the solution and care to prevent foaming. Mixing of the solution is continued for not less than 30 minutes at 3 to  $11^\circ\text{C}$  to obtain a Second Glycine Precipitate. The Second Glycine Precipitate is removed from the suspension by centrifugation or filtration and may be held frozen prior to further processing.

The Second Glycine Precipitate is solubilized in about  $9 \pm 2$  kg of citrate saline buffer ( $0.02 \pm 0.005$  M sodium citrate;  $0.12 \pm 0.03\text{M}$  sodium chloride) per kg of precipitate by mixing continuously at  $30 \pm 4^\circ\text{C}$ . The Second Glycine Precipitate solution is kept at  $30 \pm 4^\circ\text{C}$ . The pH of the Second Glycine Precipitate solution is adjusted to  $6.8 \pm 0.3$  if necessary with 1 N acetic acid or 1 N sodium hydroxide.

The procedure for preparing the Second Glycine Precipitate is repeated to obtain a Third Glycine Precipitate. The Third Glycine Precipitate solution is clarified by centrifugation and/or filtration at a temperature of  $27 \pm 7^\circ\text{C}$ .

### Example 2

Fibrinogen preparations were prepared generally as set forth in Example 1. 10 kg of the Third Glycine Precipitate were mixed with a 4:1 ratio of buffer containing 0.02M sodium citrate,

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0.12M sodium chloride, 3.2% sucrose, pH 6.7 at 30°C until in solution. Insoluble material was removed by centrifugation. The solution was diafiltered against citrate-saline buffer and concentrated to about 3% fibrinogen. The solution was sterile filtered, filled into vials (50 mL in 100 mL vials), freeze-dried and stored in the lyophilized state. A number of vials of the lyophilized preparation were reconstituted with 50 mL of water and the contents pooled. The pooled material was diafiltered against a standard formulation buffer containing 8 mM sodium citrate, 50 mM Tris, 80 mM NaCl, 50 mM glycine, pH 6.7. The solution was concentrated to about 2.3% fibrinogen and aliquoted. Appropriate amounts of stock excipient solutions were added to each aliquot to obtain the final excipient concentrations shown in Table I. The solutions containing added excipients were filled into vials (8 mL in a 20 mL vial) and lyophilized. Following lyophilization, some of the vials were heated at 80°C for 72 hours to inactivate viruses, and the vial contents were then reconstituted with 2 mL of water, where the reconstitution times and fibrinogen concentrations are set forth in Table I below.

**TABLE I**  
**RECONSTITUTION TIMES (MINUTES) OF FIBRINOGEN PRODUCT**  
**FORMULATED WITH VARIOUS EXCIPIENTS IN STANDARD BUFFER**

EXCIPIENT	Fill Conc (%)	Ex. 2A 5% Protein	Ex. 2B 6.5% Protein
No excipient		>30,>30	>30
Tween 80	0.01		9,13,5
	0.005		43,24
Tween 80/Sucrose	0.01/1		9,8
	0.005/1		21,22
Tween 80/Lys	0.01/1		6,3,6,5
	0.005/1		12,13,14
Tween 80/Lys/Sucrose	0.01/1/1		12,11
	0.005/1/1		27,15
HP $\beta$ CD (Cerestar)	1.0	4,4	10,9
HP $\alpha$ CD	1.0	2,2	10,12
HP $\gamma$ CD	1	1,4	27,16
Hydroxyethyl $\beta$ CD	1	2,4	14,16
Hydroxyethyl $\alpha$ CD	1	10,4	18,20
CarboxyMethyl $\beta$ CD	1	4,6	15,14
Methyl $\beta$ CD	1	25,4	17,33
Quaternary amine $\beta$ CD	1	24,15	26,28
Quaternary amine $\gamma$ CD	1	26,15	27,36
Tertiary amine $\beta$ CD	1	6,4	>30,18,35
Tertiary amine CarboxyMethyl $\beta$ CD	1	4,7	>30,18,19

**Example 3**

A Third Glycine Precipitate fibrinogen preparation was prepared as described in Example 1. Portions of precipitates were resuspended with a 6:1 ratio of buffer containing 20 mM citrate and 124 mM sodium chloride, pH 6.7 at 30°C until in solution. The solution was diafiltered against standard formulation buffer and concentrated to about 3% fibrinogen, and appropriate amounts of stock excipient solutions were added to obtain the final excipient concentrations shown in Tables IIA and IIB, below. The solutions containing added excipients were filled into vials (9.0 to 15.0 mL in a 20 mL vial) and lyophilized. Following lyophilization, some of the vials were heated at 80°C for 72 hours to inactivate viruses, and the vial contents were then reconstituted with 1/3 to 1/2 the fill volume of water, where the reconstitution times and protein concentrations are set forth in Tables IIA and IIB. Other vials were also reconstituted with water at 1/3 to 1/2 the original fill volume without heating, where the reconstitution times and fibrinogen concentrations are set forth in Table III below.

**TABLE IIA**  
**RECONSTITUTION TIMES (MINUTES) OF FIBRINOGEN PRODUCT**  
**FORMULATED WITH VARIOUS EXCIPIENTS IN STANDARD BUFFER**

EXCIPIENT	Fill Conc (%)	Ex. 3A 6.5% Protein	Ex. 3B 7.4% Protein
No excipient		>40	Not Done
Tween 80	0.04		>30,>30,29,28
	0.02	9,9,15,12,8,14,25,16	
	0.01	18,20,25,19,>30,>30	
Tween 80/Lysine	0.04/1	7,7,4,5	14,18,10
	0.02/1	15,21,9	>30,>30
Albumin	1.5	>35,43,48	
	1.0	33,18,27	
HP $\beta$ CD (Cerestar)	1.5	10,13,4	
HP $\beta$ CD/Tween 80	1/.02		24,23
HP $\alpha$ CD	1.5	13,5,14,13	
HP $\alpha$ CD/Tween 80	1.5/2.0		18,18
HP $\gamma$ CD	1.5	26,20	
HP $\gamma$ CD/Tween 80	1.5/.02		22,25,20
Hydroxyethyl $\beta$ CD	1.5	28, >30	
Hydroxyethyl $\alpha$ CD	1.5	>30,>30	
CarboxyMethyl $\beta$ CD	1.5	12,12,29,18,19	
CarboxyMethyl $\beta$ CD/ Tween 80	1.5/.02		21,15,17



**TABLE IIB**  
**RECONSTITUTION TIMES (MINUTES) OF FIBRINOGEN PRODUCT FORMULATED**  
**WITH VARIOUS EXCIPIENTS IN STANDARD BUFFER**

EXCIPIENT	Fill conc (%)	Ex. 3C 7% Protein	Ex. 3D 6.5% Protein	Ex. 3E 7% Protein	Ex. 3F 7.5% Protein
No excipient			>100, >100		>18 hours
Tween 80/Lys	0.16/1 0.08/1 0.04/1	40,36,39,15, 17,22, 42,20	4,6,14,7 4,10,23,12	42,18,31,40	24,26,24,21
Albumin/ Tween 80	1.0/.16			>24 hours	
HP $\beta$ CD	3.0 (Janssen) 2.5 (Cerestar) 2.0 (Janssen)		21,32,35 24,43,32	38,20,24,37,40	18,21,9,16,25
HP $\beta$ CD	1.5 (Janssen) 1.5 (Cerestar)	13,11,12,13, 13,12,7,8	20,20,31,19 12,14,11		
HP $\alpha$ CD	4 2.5 1.5	35,29,34,27, 31,24,16,30, 21	26,37,29,32	11,9,10,14,15 44,28,38,30 45,50	14,17,13,17,11 12,18,10,16,17
HP $\alpha$ CD/ Tween 80	2.5/0.16			22,35,39,23	
HP $\alpha$ CD/Lys	4.0/1 2.5/1			8,9,10,7,8	7,4,4,10,10, 16,13,10,14,10, 10
HP $\alpha$ CD/Lys/ Tween 80	4.0/1/0.16				7,8,10,12

**TABLE IIB (Continued)**  
**RECONSTITUTION TIMES (MINUTES) OF FIBRINOGEN PRODUCT FORMULATED**  
**WITH VARIOUS EXCIPIENTS IN STANDARD BUFFER**

<b>EXCIPIENT</b>	<b>Fill conc (%)</b>	<b>Ex. 3C 7% Protein</b>	<b>Ex. 3D 6.5% Protein</b>	<b>Ex. 3E 7% Protein</b>	<b>Ex. 3F 7.5% Protein</b>
HP $\gamma$ CD	4			87,110	
Hydroxyethyl $\alpha$ CD	4			41,68	
CarboxyMethyl $\beta$ CD	4			97,46,35	
Tertiary amine $\beta$ CD	4			67	
Tertiary amine CarboxyMethyl $\beta$ CD	4			99	

**Table III**  
**Fibrinogen Formulations and Reconstitution Times**  
**for Non Heat-Treated Fibrinogen Vials**

	Excipient	Excipient Conc.	Fill Vol. (mL)	Reconst. Vol. (mL)	Fibrin. (mg/mL)	Reconst. Time
Ex. 3G	HP $\alpha$ CD	1.5%	12.2	5	ND	16 min.
	HP $\alpha$ CD	1.5%	10.6	5	ND	17 min.
	HP $\beta$ CD	1.5%	12.2	5	ND	10 min.
	HP $\beta$ CD	1.5%	10.6	5	ND	3 min.
Ex. 3H	HP $\alpha$ CD	1.5%	13	5	67	17, 19 min.
	HP $\beta$ CD	1.5%	13	5	64	11, 12 min.
	No Excipient	N/A	13	5	66	> 24 h, > 24 h
Ex. 3I	HP $\alpha$ CD	1.5%	12.5	5	64	39 min.
	HP $\alpha$ CD	2.5%	12.5	5	69	23, 33 min.
	HP $\alpha$ CD	4%	12.5	5	73	11.5, 13 min.
	HP $\beta$ CD	2.5%	12.5	5	69	21, 7 min.
	No Excipient	N/A	12.5	5	ND	> 7.5 h
Ex. 3J	HP $\alpha$ CD	2.5%	15	5	76	11, 15 min.
	HP $\alpha$ CD	4%	15	5	77	11.5, 15 min.
	HP $\beta$ CD	2.5%	15	5	79	11, 11 min.
	No Excipient	N/A	15	5	81	> 24 h, > 24 h

N/A = Not Applicable; ND = Not Determined

#### Example 4

A Third Glycine Precipitate fibrinogen preparation was prepared as described in Example 1. Portions of precipitates were resuspended with a 6:1 ratio of buffer containing 20 mM citrate and 124 mM sodium chloride, pH 6.7 at 30°C until in solution. The solution was diafiltered against standard formulation buffer and concentrated to about 3% fibrinogen, and appropriate amounts of stock excipient solutions were added to obtain the final excipient concentrations shown in Table IV, below. The solutions containing added excipients were filled into vials (15.0 mL in a 20 mL vial) and lyophilized. Following lyophilization, some of the vials were heated at 80°C for 72 hours to inactivate viruses, and the vial contents were then reconstituted with water at  $\frac{1}{3}$  to  $\frac{1}{2}$  the original fill volume, where the reconstitution times and protein concentrations are set forth in Table IV. Other vials were not heat treated and vial

contents were reconstituted with 5 mL of water, where the reconstitution times and fibrinogen concentrations are set forth in Table V.

**TABLE IV**  
**RECONSTITUTION TIMES (MINUTES) AFTER HEAT TREATMENT OF FIBRINOGEN**  
**PRODUCT FORMULATED WITH VARIOUS EXCIPIENTS IN STANDARD BUFFER\***

EXCIPIENT	Fill Conc (%)	Ex. 4A 7.4% Protein	Ex. 4B 6.4% Protein	Ex. 4C 8.6% Protein	Ex. 4D	Ex. 4E
No excipient		>30,>30				
HP $\alpha$ CD	4.0	10,6,6	5,5,6,6,2	7,6,6	8,8,7	10,10
	2.5	7,5,9	5,6,7,5,5			
HP $\alpha$ CD/Lys	4.0/1	3,4,4,2,2,5	4,4,6,6	4,4,5	5,2	
	2.5/1	5,5,7,3,4,3	5,5,2,8,6			
Lys	1.0	>30,>30,>30				
HP $\beta$ CD	2.5	7,6,8	14,12			
	1.5	5,9,17	11,10,11			

\* All vials were filled with 13.6 to 15 mL of product and reconstituted with 5 mL of sterile water.

<b>Table V</b> <b>Fibrinogen Formulations and Reconstitution Times</b> <b>for Non Heat-Treated Fibrinogen Vials</b>						
	<b>Excipient</b>	<b>Excipient Concen.</b>	<b>Fill Vol. (mL)</b>	<b>Reconst. Vol. (mL)</b>	<b>Fibrin. (mg/mL)</b>	<b>Reconst. Time</b>
Ex. 4F	HP $\alpha$ CD	2.5%	15	5	84	10, 6 min.
	HP $\alpha$ CD	4%	15	5	82	3, 4 min.
	HP $\beta$ CD	2.5%	15	5	82	4.5, 7 min.
	HP $\beta$ CD	1.5%	15	5	84	8.5, 12 min.
	No Excipient	N/A	15	5	83	> 60, > 80 min.
Ex. 4G	HP $\alpha$ CD	2.5%	15	5	69	5.5, 7, 6, 9 min.
	HP $\alpha$ CD	4%	15	5	63	4, 6, 4, 5, 5 min.
	HP $\beta$ CD	2.5%	15	5	66	10, 7 min.
	HP $\beta$ CD	1.5%	15	5	66	7.5, 9 min.
	No Excipient	N/A	15	5	ND	> 2 h, > 4 h
Ex. 4H	HP $\alpha$ CD	4%	15	5	ND	4, 7 min.
Ex. 4I	HP $\alpha$ CD	4%	15	5	ND	9, 6 min.
Ex. 4J	HP $\alpha$ CD	4%	15	5	ND	15 min.

N/A = Not Applicable

ND = Not Determined

The above descriptions of exemplary embodiments of processes for preparing stabilized fibrinogen products are for illustrative purposes. Because of variations that will be apparent to those skilled in the art, the present invention is not intended to be limited to the particular embodiments described above. This invention can also be practiced in the absence of any element not specifically disclosed. The scope of the invention is described in the following claims.